

Letter: Desiccated Thyroid Extract Causes Nonphysiologic T₃ Peaks

To the Editor:

Hoang and associates once again raise the old question of a possible place for animal sources of desiccated thyroid extract (DTE) in the treatment of hypothyroidism (1). They stimulate new discussion but do not seem to break new ground. Of the 70 patients in their 16-week crossover study comparing DTE with l-thyroxine (L-T₄), 49% preferred DTE and reported a subjective improvement in quality of life, although there were no statistically significant differences between the groups on psychometric testing. Those preferring DTE also had a 3- to 4-lb weight loss, but started with higher mean body weights than those in the L-T₄ group (178.95 lb vs. 162.80 lb). Thyroid biochemical tests were within normal ranges in both groups. However, serum T₃ levels were statistically higher (P<0.001) in the subjects who preferred DTE when measured just before the once-daily dose of DTE (at the expected daily nadir). In the only 2 subjects given DTE who had T₃ measured both before and again 3 hours after DTE administration (near the expected T₃ peak levels) serum T₃ rose 23% and 36%. Such nonphysiologic changes in serum T₃ after DTE administration and resultant risks have long been known (2) and are the subject of concern (3). While Hoang et al. do briefly mention possible cardiovascular risks from these T₃ changes in the body of their report and in their supplementary data, they do not comment on such risks in their abstract, nor do they discuss potential adverse effects on bone turnover at all.

Exploring a role for DTE in the treatment of hypothyroidism with a well-designed, blinded, randomized clinical trial is laudable. However, when evaluating a therapy for a condition that affects millions of patients and for which an effective treatment already exists (4), this clinical trial should be powered and designed to detect adverse consequences. When the goal is physiologic replacement, care also needs to be exercised that normal physiology is restored. The study of Hoang and colleagues is provocative, but it does not achieve the minimum standard required to alter current clinical practice.

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References

1. Hoang TD, Olsen CH, Mai VQ, Clyde PW, Shakir MK. Desiccated thyroid extract compared with levothyroxine in the treatment of hypothyroidism: a randomized, double blind, crossover study. *J Clin Endocrinol Metab* 2013;98:1982-90. Epub March 28, 2013.
2. Saberi M, Utiger RD. Serum thyroid hormones and thyrotropin concentrations during thyroxine and triiodothyronine therapy. *J Clin Endocrinol Metab* 1974;39:923-7.
3. Biondi B, Wartofsky L. Combination treatment with T₄ And T₃: toward personalized replacement therapy in hypothyroidism? *J Clin Endocrinol Metab* 2012;97:2256-71. Epub May 16, 2012.
4. Garber JR, Cobin RJ, Gharib H, et al. Clinical practice guidelines for hypothyroidism in adults: cosponsored by the American Association of Clinical Endocrinologists and the American Thyroid Association. *Thyroid* 2012; 22:1200-35. Epub November 6, 2012.